



Billing Code 4410-09-M

DEPARTMENT OF JUSTICE  
DRUG ENFORCEMENT ADMINISTRATION  
IMPORTER OF CONTROLLED SUBSTANCES  
NOTICE OF REGISTRATION  
MYLAN PHARMACEUTICALS INC.

By Notice dated December 22, 2011, and published in the  
Federal Register on December 29, 2011, 76 FR 81978, Mylan  
Pharmaceuticals, Inc., 781 Chestnut Ridge Road, Morgantown, West  
Virginia 26505, made application by renewal to the Drug  
Enforcement Administration (DEA) to be registered as an importer  
of the following basic classes of controlled substances:

Drug	Schedule
Methylphenidate (1724)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Fentanyl (9801)	II

The company plans to import the listed controlled  
substances in finished dosage form (FDF) from foreign sources  
for analytical testing and clinical trials in which the foreign  
FDF will be compared to the company's own domestically-  
manufactured FDF. This analysis is required to allow the

company to export domestically-manufactured FDF to foreign markets.

No comments or objections have been received. DEA has considered the factors in 21 USC § 823(a) and § 952(a), and determined that the registration of Mylan Pharmaceuticals, Inc. to import the basic classes of controlled substances is consistent with the public interest, and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. DEA has investigated Mylan Pharmaceuticals, Inc. to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 USC § 952(a) and § 958(a), and in accordance with 21 CFR § 1301.34, the above named company is granted registration as an importer of the basic classes of controlled substances listed.

Joseph T. Rannazzisi  
Deputy Assistant Administrator  
Office of Diversion Control  
Drug Enforcement Administration

DATED: February 23, 2012

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